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54 Room-temperature-curable organopolysiloxane composition.

57 A room-temperature-curable organopolysiloxane composition having a highly durable adhesion that bonds well to various substrates, e.g. glass, plastics, and metals, through contact-curing and exhibits an excellent early-cure adhesiveness and that cures to give a silicone rubber that is highly water-resistant and exhibits long-term retention of its bonding strength even in hostile ambients, e.g. immersion in hot water, comprises a composition of a hydroxyl-terminated or alkoxy-terminated organopolysiloxane, an inorganic filler, an rosin ester in which the ester is a polyol, a reaction mixture from aminoalkylalkoxysilane and epoxyalkylalkoxysilane, and a curing catalyst.

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The present invention relates to a room-temperature-curable organopolysiloxane composition. More specifically, the present invention relates to a room-temperature-curable organopolysiloxane composition that develops a strong adhesion for a variety of substrates, e.g., glass, plastics, metals, and so forth, when cured while in contact with the substrate (contact-curing) and, moreover, that develops this adhesion at an early stage in its cure, wherein this adhesion is highly durable to challenge by hostile ambients, e.g., immersion in hot water and so forth.

Mayuzumi et al in U.S. Patent No. 3,837,876, issued September 24, 1974, and Joseph et al in U.S. Patent No. 4,602,078, issued July 22, 1986, teach organopolysiloxane compositions that are adhesive for a variety of substrates through contact-curing. These compositions consist of hydroxyl-endblocked or-ganopolysiloxane, aminoalkylalkoxysilane/epoxyalkylalkoxysilane mixture or reaction product, and curing catalyst. However, the adherence of the silicone rubbers afforded by the cure of these compositions deteriorates upon exposure to water. In particular, their bonding strength for float glass deteriorates during challenge by hostile ambients, such as immersion in hot water and the like.

The present inventors carried out extensive investigations directed at solving the above-described problems. As a result, they discovered that a highly durable adhesion could be obtained from a room-temperature-curable organopolysiloxane composition that contained an aminoalkylalkoxysilane/epoxyalkylalkoxysilane reaction mixture and resin acid including rosin. However, it was found that during the early stage of its cure this composition is slow to develop adhesive strength for certain substrates. Accordingly, the inventors continued their investigations and were able to achieve the present invention as a result.

The present invention takes as its object the introduction of a room-temperature-curable or-ganopolysiloxane composition having a highly durable adhesion that develops an excellent early-cure adhesiveness for a variety of substrates, e.g., glass, plastics, and metals, by contact-curing and that cures to afford a highly water-resistant silicone rubber that manifests a long-term retention of its bonding strength even when challenged by a hostile ambient, e.g., hot-water immersion.

This invention relates to a room-temperature-curable organopolysiloxane composition comprising (A) 100 parts by weight of hydroxyl-endblocked or alkoxy-endblocked organopolysiloxane that has a viscosity at 25°C within the range of 0.000020 to 1 m<sup>2</sup>/s, (B) 1 to 300 parts by weight inorganic filler, (C) 0.1 to 10 parts by weight rosin ester which is an esterification product of rosin and a polyol, (D) 0.5 to 30 parts by weight reaction mixture from aminoalkylalkoxysilane and epoxyalkylalkoxysilane, and (E) 0.001 to 10 parts by weight curing catalyst.

The organopolysiloxane of the component (A) used in the present invention is the main or base component of the composition. The organopolysiloxane must be endblocked by hydroxyl groups or alkoxy groups. When its viscosity is too low, the post-cure rubber elasticity will be poor. On the other hand, the processability is impaired when its viscosity is too high. For these reasons, its viscosity at 25°C must fall within the range of 0.000020 to 1 m<sup>2</sup>/s (square meters per second), while the preferred viscosity range is 0.0001 to 0.1 m<sup>2</sup>/s. This organopolysiloxane preferably has a straight-chain molecular structure, but a moderately branched chain is also usable.

The organic groups in this organopolysiloxane are exemplified by alkyl groups such as methyl, ethyl, propyl, butyl, and octyl; alkenyl groups such as vinyl, allyl, and hexenyl; aryl groups such as phenyl and tolyl; and substituted alkyl groups such as 3,3,3-trifluoropropyl, 3-chloropropyl, and 3-cyanoalkyl. The organopolysiloxane under consideration is specifically exemplified by dimethylpolysiloxanes, methylethylpolysiloxanes, methyloctylpolysiloxanes, methylvinylpoly-siloxanes, methylphenylpolysiloxanes, methyl-(3,3,3-trifluoro-propyl)polysiloxanes, dimethylsiloxane-methylphenylsiloxane copolymers, and dimethylsiloxane-methyl(3,3,3-trifluoropropyl)-siloxane copolymers. The hydroxyl-endblocked molecular chain terminals are preferably dimethylhydroxysiloxy and methylphenylhydroxysiloxy. The alkoxy-endblocked molecular chain terminals are preferably by vinyl dimethoxysiloxy, methyl dimethoxysiloxy, trimethoxysiloxy, methyldiethoxysiloxy, and triethoxysiloxy.

The inorganic filler of the component (B) employed by the present invention functions to thicken the composition and to provide the cured product therefrom with mechanical strength. This inorganic filler is exemplified by alkaline-earth metal salts, inorganic oxides, metal hydroxides, and carbon black. The alkaline-earth metal salts are specifically exemplified by the carbonates, bicarbonates, sulfates, etc., of calcium, magnesium, and barium. The inorganic oxides are specifically exemplified by fumed silica, calcined silica, precipitated silica, microparticulate quartz, titanium oxide, diatomaceous earth, alumina, and the like. The metal hydroxides are specifically exemplified by aluminum hydroxide. These fillers may also be used after surface treatment with a silane, silazane, low degree-of-polymerization siloxane, or an organic compound.

Among the preceding inorganic fillers, fumed silica, precipitated silica, and colloidal calcium carbonate are preferred when higher mechanical strengths are desired for the cured product.

Component (B) is added at 1 to 300 parts by weight per 100 parts by weight component (A). Component (B) may take the form of a single species of inorganic filler or any desired mixture of the preceding. The cured product obtained from the composition under consideration will have a poor mechanical strength at a component (B) addition of less than 1 part by weight. On the other hand, when the component (B) addition exceeds 300 parts by weight, it becomes highly problematic to produce a cured product having a good rubber elasticity.

The rosin ester of the component (C) employed by the present invention, through its joint use with component (D), functions to provide the cured product obtain from the inventive composition with adhesiveness in hostile ambients, e.g. immersion in hot water, without impairing the early - cure development of adhesiveness. This rosin ester is the product obtained by the alcohol esterification of rosin. Rosin is the turpentine oil-free nonvolatile component obtained from the steam distillation of the resin from coniferous trees, such as pine and so forth. The alcohol used in this esterification is a polyol having two or more C-OH groups per molecule, for example, ethylene glycol, glycerol, and pentaerythritol. Although rosin typically contains 2 double bonds, there is no impediment to the use of ester compounds deriving from modified rosin that has been modified by disproportionation, dimerization, or hydrogenation.

Component (C) should be added at 0.1 to 10 parts by weight per 100 parts by weight component (A), preferably from 0.5 to 5 parts by weight. A satisfactorily durable adhesion is not developed at an addition below 0.1 weight parts, while additions in excess of 10 parts by weight cause a slow curing rate and a diminished post-cure rubber strength.

Component (D) comprises the reaction mixture from aminoalkylalkoxysilane and epoxyalkylalkoxysilane. This component functions as a crosslinker for the composition and also serves to impart the capacity to adhere to various types of substrates by contact-curing. In addition, component (D), through its joint use with component (C), provides the cured product from the invention composition with an adherence that is durable to hostile ambients, e.g. immersion in hot water.

The aminoalkylalkoxysilane making up component (D) is exemplified by aminomethyltriethoxysilane, gamma-aminopropyl-triethoxysilane, gamma-aminopropylmethyldimethoxysilane, N-(beta-aminoethyl)aminomethyltributoxysilane, N-(beta-aminoethyl)-gamma-aminopropyltrimethoxysilane, N-(beta-aminoethyl)-gamma-aminopropylmethyldimethoxysilane, and gamma-anilinopropyl-triethoxysilane.

The epoxyalkylalkoxysilane making up this component (D) is exemplified by gamma-glycidoxypropyl-trimethoxysilane, gamma-glycidoxypropylmethyldimethoxysilane, beta-(3,4-epoxycyclohexyl)-ethyltrimethoxysilane, and beta-(3,4-epoxycyclohexyl)ethylmethyldimethoxysilane.

The reaction mixture is prepared simply by mixing the aminoalkylalkoxysilane and epoxyalkylalkoxysilane at a molar ratio in the range of (1:1.5) to (1:5) and preferably (1:2) to (1:4) and heating or holding at room temperature.

Component (D) should be added at 0.5 to 30 parts by weight per 100 parts by weight component (A) and is preferably added at 1 to 10 parts by weight per 100 parts by weight component (A). The basis for this range is as follows: an adequate rubber strength and adhesiveness are not obtained when less component (D) is present; on the other hand, when more is present, the curing rate is diminished and the cured rubber is too hard.

The curing catalyst of component (E) functions as a catalyst to accelerate the cure of the composition according to the present invention. Component (E) is exemplified by the carboxylic acid salts of tin, titanium, zirconium, iron, antimony, bismuth, and manganese, and by organotitanate esters and organotitanium chelate compounds. Operative catalysts are specifically exemplified by tin compounds such as dibutyltin dilaurate, dibutyltin dioctate, dioctyltin dilaurate, dibutyltin malate ester, and stannous octoate, and by titanium compounds such as tetrabutyl titanate, diisopropoxybis(acetyl-acetate)titanium, and diisopropoxybis(ethyl acetoacetate)-titanium.

Component (E) should be added at 0.001 to 10 parts by weight per 100 parts by weight component (A) and is preferably added at 0.01 to 5 parts by weight per 100 parts by weight component (A). The basis for this range is as follows: when less than 0.001 part by weight of component (E) is present, the curing rate is so slow that practical use is precluded; the presence of more than 10 parts by weight of component (E) causes such a high curing rate that the use time is eliminated.

In order to adjust the curability and post-cure rubber strength of the composition according to the present invention, it may additionally contain (F) an alkoxysilane crosslinker different from component (D). This alkoxysilane is exemplified by tetrafunctional alkoxysilanes such as tetramethoxysilane, tetraethoxysilane, methyl Cellosolve orthosilicate, and n-propyl orthosilicate; by trifunctional alkoxysilanes such as

methyltri - methoxysilane, methyltriethoxysilane, ethyltrimethoxysilane, vinyltrimethoxysilane, phenyl - trimethoxysilane, and methyltri - (methoxyethoxy)silane; and by the partial hydrolyzates of the preceding. The alkoxysilane crosslinker is preferably added in an amount of from 1 to 10 parts by weight based on 100 parts by weight of (A).

In addition, the composition according to the present invention may contain the following on an optional basis insofar as the object of the present invention is not impaired: organic solvent, trimethylsiloxy - terminated diorganopolysiloxane, flame retardant, plasticizer, thixotropy agent, colorant, ordinary adhesion promoters, and antimolds.

The present invention is explained in greater detail below through illustrative examples. In the examples and comparison examples, parts designates parts by weight, the viscosity is the value at 25 ° C, and the following abbreviations are used: Tmax = maximum tensile stress; Emax = elongation at maximum load; and CF ratio = cohesive failure ratio in the sample.

#### EXAMPLE 1

The following were mixed to homogeneity to afford a mixture hereinafter designated as the "base": 100 parts hydroxyl - endblocked polydimethylsiloxane with viscosity = 0.015 m<sup>2</sup>/s, 100 parts fatty acid - treated calcium carbonate (average particle diameter = 0.08 micrometers), and 2 parts ethylene glycol ester of rosin (viscosity = 200 Pa.s).

The following were also mixed to afford a mixture hereinafter designated as the "catalyst": 60 parts n - propyl orthosilicate, 40 parts of the 1:2 molar ratio reaction mixture from gamma - aminopropyltriethoxysilane and gamma - glycidoxypropyl - trimethoxysilane, and 1 part dibutyltin dilaurate.

The base and catalyst were then mixed at a 100:5 weight ratio to give a room - temperature - curable organopoly - siloxane composition. Tensile adhesion test specimens were prepared based on JIS A 5758 using this composition. The adherends were float glass and high - performance heat - reflecting glass (SGY32 from Asahi Glass Kabushiki Kaisha). The tensile adhesion was measured after curing for 5 days at 20 ° C. In addition, the tensile adhesion was also measured after curing for 7 days at 20 ° C and then 7 days at 50 ° C and then immersion for 1 month in hot water (80 ° C). These measurement results are reported in the Table, respectively, in the columns labeled "after cure for 5 days at 20 ° C" and "post - immersion in 80 ° C water".

#### EXAMPLE 2

A room - temperature - curable organopolysiloxane composition was prepared as in Example 1, but in the present case using 2.5 parts of a toluene solution (solids = 80%) of glycerol ester of rosin (ester softening point = 90 ° C) in place of the ethylene glycol ester of rosin used in Example 1. This composition was evaluated as in Example 1, and these results were as reported in the Table.

#### EXAMPLE 3

A room - temperature - curable organopolysiloxane composition was prepared as in Example 1, but in the present case using 2.5 parts of a toluene solution (solids = 80%) of pentaerythritol ester of rosin (ester softening point = 120 ° C) in place of the ethylene glycol ester of rosin used in Example 1. This composition was evaluated as in Example 1, and these results were as reported in the Table.

#### EXAMPLE 4

A room - temperature - curable organopolysiloxane composition was prepared as in Example 1, but in the present case using 2.5 parts of a toluene solution (solids = 80%) of glycerol ester of hydrogenated rosin (ester softening point = 90 ° C) in place of the ethylene glycol ester of rosin used in Example 1. This composition was evaluated as in Example 1, and these results were as reported in the Table.

#### COMPARISON EXAMPLE 1

A room - temperature - curable organopolysiloxane composition was prepared as in Example 1, but in the present case using 2.5 parts of a toluene solution of rosin (solids = 80%) in place of the ethylene glycol ester of rosin used in Example 1. This composition was evaluated as in Example 1, and these results were as reported in the Table.

## COMPARISON EXAMPLE 2

A room-temperature-curable organopolysiloxane composition was prepared as in Example 1, but in the present case omitting the ethylene glycol ester of rosin from the base of Example 1. This composition was evaluated as in Example 1, and these results were as reported in the Table.

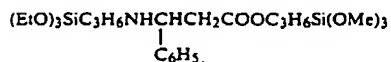
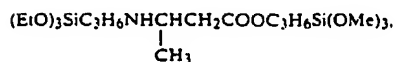
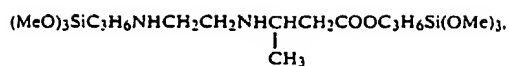
TABLE

Composition Number	Adherend	After cure for 5 days at 20°C				Post-immersion in 80°C water		
		Tmax 2 kgf/cm <sup>2</sup>	Emax %	CF ratio %		Tmax 2 kgf/cm <sup>2</sup>	Emax %	CF ratio %
Example 1	float glass	10.7	170	100		10.3	150	100
	SGY32	10.6	160	100		11.1	160	100
Example 2	float glass	10.2	160	100		10.9	150	100
	SGY32	10.0	160	100		10.5	140	100
Example 3	float glass	11.2	160	100		11.4	150	100
	SGY32	10.5	150	100		10.4	140	100
Example 4	float glass	10.9	160	100		10.3	140	100
	SGY32	10.4	140	100		11.3	160	100
Comparison Example 1	float glass	10.3	150	100		10.7	140	100
	SGY32	5.6	50	10		10.6	150	100
Comparison Example 2	float glass	10.0	160	100		6.6	90	40
	SGY32	10.5	160	100		10.2	140	100

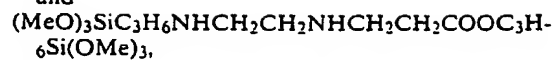
The room - temperature - curable organopolysiloxane composition according to the present invention characteristically bonds well to substrates through contact - curing, exhibits an excellent early - cure adhesion, and after curing is highly water - resistant and exhibits long - term retention of its bonding strength even in hostile ambients e.g., immersion in hot water, because it is composed of components (A) through (E) and in particular because it contains the rosin ester of a polyol as component (C) and the aminoalkylalkoxysilane/epoxyalkylalkoxysilane reaction mixture of component (D).

# Claims

1. A room - temperature - curable organopolysiloxane composition comprising  
(A) 100 parts by weight of hydroxyl - endblocked or alkoxy - endblocked organopolysiloxane that has a viscosity at 25 ° C within the range of 0.000020 to 1 m<sup>2</sup>/s,  
(B) 1 to 300 parts by weight inorganic filler,  
(C) 0.1 to 10 parts by weight rosin ester which is an esterification product of rosin and a polyol,  
(D) 0.5 to 30 parts by weight reaction mixture from aminoalkylalkoxysilane and epoxyalkylalkoxysilane, and  
(E) 0.001 to 10 weight parts curing catalyst.
2. The room - temperature - curable organopolysiloxane composition according to claim 1 further comprising (F) an alkoxysilane crosslinker.



and



wherein Me is the methyl group and Et is the ethyl group.

5 5. The composition according to claim 1, wherein the organosilicon compound (C) is at least one selected from the group consisting of alkoxysilane, ketoximesilane, acyloxysilane, alkenyloxysilane, amidosilane, 10 aminosilane, aminoxysilane and partially hydrolyzed products of these silanes.

6. A cured product obtained by curing the composition as claimed in claim 1.

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## EUROPEAN SEARCH REPORT

Application Number

EP 92 11 8641

DOCUMENTS CONSIDERED TO BE RELEVANT			
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int. Cl.5)
Y	DATABASE WPIL Section Ch, Week 8931, 27 September 1989 Derwent Publications Ltd., London, GB; Class A81, AN 89-225725 & JP-A-1 163 281 (YOKOHAMA RUBBER KK) * abstract *	1,2	C08L83/04
Y	US-A-3 542 714 (N. T. METTERS) * claims 1,2; examples 1-3 * * column 2, line 19 - line 54 *	1,2	
A	CHEMICAL ABSTRACTS, vol. 115, no. 2, 15 July 1991, Columbus, Ohio, US; abstract no. 11055r, FUJIMOTO, T. ET AL. 'Room-temperature-curable silicone rubber compositions' page 125 ;column L ; * abstract * & JP-A-0 317 158 (TOSHIBA SILICONE CO., LTD.)	1,2	
A	GB-A-2 065 152 (IMPERIAL CHEMICAL INDUSTRIES LTD.) * page 2, line 1 - line 9; claims 1-3 *	1	TECHNICAL FIELDS SEARCHED (Int. Cl.5)
D,A	US-A-4 602 078 (E. A. JOSEPH ET AL.) * column 4, line 57 - line 64; claims 1-5 *	1,2	C08L C09D
The present search report has been drawn up for all claims			
Place of search THE HAGUE		Date of completion of the search 09 DECEMBER 1992	Examiner KANETAKIS I.
<b>CATEGORY OF CITED DOCUMENTS</b> X : particularly relevant if taken alone Y : particularly relevant if combined with another document of the same category A : technological background O : non-written disclosure P : intermediate document T : theory or principle underlying the invention E : earlier patent document, but published on, or after the filing date D : document cited in the application L : document cited for other reasons & : member of the same patent family, corresponding document			

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